



(10) International Publication Number
WO 03/080646 A2

PCT

(84) Designated States (regional): ARIPO patent (GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, BG, CH, CY, CZ, DE, DK, EE,

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1 GAATTCTCTCCAGCTCAGCAGCCGCCCGCCAGAGCAGAGCAACCCGCAATCCGACGACATC 60
 CTTAAGCAGCTCGAATGTCCTGGCGCCGGTCTCTGCTCTGCTTGGCGGTTAGGCTTCGGTGG
 61 TCTGGAACCTTCAGATGTCAGATATGTCCTCAGCCCTCACTGCTAGTCTCTGGGCGCTGGCC
 120 AGACTCTTGAAGTCTCTAGCTCTTACAGAGTTCGGAGTGGAGCGATCAGGACCCGCGACCGG
 aa I Q N S P A L T C L V L G L A -
 [signal Peptide]
 121 CTGTGCTTTGGTGAAGGGTCTGCTGTGTGCACCACTCCCCCATCTACTGTGGCCCACTGGCC
 180 GAACAGAAACCCTCTCCAGACACACCTGTGGAGGGGTAGAGATCAGCGGGTGGACCGG
 aa L V F F G E G S A I V E H F P S Y V A N L A I
 Start Mature Protein
 181 TCAGACTCTGGGTCAGGGTATTCAGCGGGTGGTGGCTTCAGGAGCTCGAAGTGG
 240 AGCTGAAAGCCCACTCCCAAAATCTGTCAACCGGTCCGGAGGTCTCTGGGGTGCAC
 aa S D F G V R V F Q Q V A Q A S K D R N V 32
 GTTTTCTACCCCTATGGGTGGCTCGGTGTTGGCCATGCTCCAGCTGACACAGGAGGA
 241 CAARAGATGGGATACCTGACCGGAGACAAAGGCTCAGAGGAGACATGTTGCTGCTCT
 aa V F S P Y G G V A S V L A M L Q L T T G G 52
 GAAACCCAGCAGCAGATTCAAGCAGCTATGGGATTCAGATGATGACRAGGSGATGGCC
 301 CTTTGGGTGCTGCTCTAAGTTCGTGATACCTAAGTCTAACTACTGTTCCGTACCGG
 aa E T Q Q Q T Q A A N G F K I D D K G M A 72
 CCCGCCCTCCGGCATCTGTACAAGGAGCTCATATGGGCCATGGAAAGGATGAGATCAGC
 361 GGGCGGAGGCCGTAGACATGTTCTCTCAGTACCCGGGTACCTTGTCTTACTCTAGTGG
 aa P A L R G L L T E K L E L N G P W N R D E I S 92
 ACCACGAGCGGATCTTCTGTCCAGCGGGATCTTAAAGTGTCCAGGGCTCATGCCCCAC
 421 TGGTGTCTGCGCTAAGAGCAGGTTCGCCCTAGACTTCGACACAGTCCCGAAGTACGGGTG
 aa T D A I F V Q R D L K L V Q G F M P E N 123
 TCTCTCAGGCTTCTCCGAGCAGCGTCAAGCAAGTGGACTTTCAGAGGTGGAGAGAGCC
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 aa F F R L F R S T V K Q V D F S E V N R A 132
 AGATTCAATCATGACTGGGTGAAGACACACACAAAGGTATGATCAGCAACTTGTCTT
 541 TCTAAGTAGTAGTACTGACCACTCTGTGTGTGTGTTTCCATCTAGTGGTTGAACGAA
 aa R F I X I D N W V K T E T K G M I S N L L 152

(57) Abstract: The present invention is based upon the discovery that modified plasminogen activator inhibitor type-I (PAI-1) in which two or more amino acid residues that do not contain a sulfhydryl group have been replaced with amino acid residues that contain a sulfhydryl group and, therefore, forms intramolecular disulfide bonds, have increased in vivo half-life. Also disclosed are the modified PAI-1 proteins, derivatives and analogs thereof, specific antibodies, nucleic acid molecules and host cells. Methods for producing modified PAI-1, derivatives and analogs are also provided. The invention further relates to Therapeutics, pharmaceutical compositions and method of using the composition for treatment. The invention may be used to inhibit angiogenesis in a subject, thereby treating diseases or conditions associated with undesired angiogenesis and cell proliferation. Such conditions include psoriasis, chronic inflammation, tumor invasion and metastasis and conditions in which angiogenesis is pathogenic. The modified PAI-1 molecules of the present invention are useful for the treatment, prophylaxis, management and amelioration of cardiovascular diseases such as, but not limited to those that are related to hyperfibrinolysis, hemophilia, and vessel leakage syndrome.